

IN THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in this application.

D | 1. (Currently Amended) A method for producing L-glutamic acid, comprising ~~the steps of~~ cultivating a coryneform bacteria in a liquid medium to produce and accumulate L-glutamic acid in the medium, and collecting the L-glutamic acid,

wherein a penicillin binding protein (PBP) is not produced or the function of a penicillin binding protein is reduced or eliminated in said coryneform bacteria due to a mutation in said produced penicillin binding protein,

wherein said penicillin binding protein is encoded by a DNA which comprises nucleotides 881 to 2623 of SEQ ID NO:1, or a DNA which is hybridizable with a nucleotide sequence comprising at least nucleotides 881 to 2623 of SEQ ID NO:1 under stringent conditions and which codes for a penicillin binding protein, wherein the stringent conditions comprise washing at 60°C in 1 X SSC and 0.1% SDS, and

wherein said bacteria have the ability to produce L-glutamic acid.

2. (Currently Amended) The method according to claim 1, wherein the ~~coryneform bacteria are bacteria in which~~ a penicillin binding protein is produced or the function of a penicillin binding protein is not reduced or eliminated at a first temperature and ~~a~~ the penicillin binding protein is not produced or the function of a penicillin binding protein is reduced or eliminated at a second temperature because of a mutation in said [produced] penicillin binding protein ~~at a second temperature~~,

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comprising ~~the steps of~~ cultivating the coryneform bacteria at the first temperature to proliferate the coryneform bacteria, and cultivating the coryneform bacteria at the second temperature to produce L-glutamic acid.

3. (Currently Amended) The method according to claim 2, wherein the coryneform bacteria ~~are bacteria which harbor~~ comprise a plasmid comprising ~~a gene~~ the DNA coding for ~~a~~ the penicillin binding protein and a temperature sensitive replication control region, and in which ~~said PBP gene~~ the DNA encoding the penicillin binding protein, which is on the a bacterial chromosome does not function, which DNA also comprises nucleotides 881 to 2623 of SEQ ID NO:1 or a DNA which is hybridizable to at least nucleotides 881 to 2623 of SEQ ID NO:1 under stringent conditions, which comprise washing at 60°C in 1 X SSC and 0.1% SDS; and the plasmid can replicate at the first temperature, and cannot replicate at the second temperature.

Claim 4 (Previously canceled).

Claim 5 (Canceled).

6. (Previously Amended) The method according to claim 1, wherein the penicillin binding protein has the amino acid sequence shown in SEQ ID NO:2.

7. (Previously Amended) The method according to claim 3, wherein the PBP gene has a nucleotide sequence comprising at least nucleotides 881 to 2623 of SEQ ID NO:1.

8. (Previously Amended) A DNA which codes for a protein which has the amino acid sequence of SEQ ID NO:2.

D2 9. (Currently Amended) A DNA derived from coryneform bacterium, said DNA is defined in the following (a) or (b):

(a) a DNA which comprises at nucleotides 881 to 2623 of SEQ ID NO:1;

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(B b) a DNA which is hybridizable with a nucleotide sequence comprising at least nucleotides 881 to 2623 of SEQ ID NO:1 under a stringent condition, which comprises washing at 60°C in 1 X SSC and 0.1% SDS, and wherein said DNA codes for a protein having the ability to bind to penicillin.

Claim 10 (Canceled)

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11. (New) The DNA of Claim 9, which is (a).

12. (New) The DNA of Claim 9, which is (b).

13. (New) A vector comprising the DNA of Claim 11.

14. (New) A vector comprising the DNA of Claim 12.

15. (New) A bacterial cell comprising the vector of Claim 13.

16. (New) A bacterial cell comprising the vector of Claim 14.

17. (New) The method according to claim 1, wherein at least a portion of the DNA which comprises nucleotides 881 to 2623 of SEQ ID NO:1 or a DNA which is hybridizable with a nucleotide sequence comprising at least nucleotides 881 to 2623 of SEQ ID NO:1 is deleted such that the function of the penicillin binding protein is reduced or eliminated.

18. (New) The method according to claim 1, wherein said penicillin binding protein is encoded by a DNA which comprises nucleotides 881 to 2623 of SEQ ID NO:1.

19. (New) The method according to claim 1, wherein said penicillin binding protein is encoded by DNA which is hybridizable with a nucleotide sequence comprising at least nucleotides 881 to 2623 of SEQ ID NO:1 under stringent conditions, which comprise washing at 60°C in 1 X SSC and 0.1% SDS.

20. (New) In a method of making a seasoning, the improvement comprising producing L-glutamic acid according to the method according to claim 1.

21. (New) In a method of making a seasoning, the improvement comprising producing L-glutamic acid according to the method according to claim 2.

22. (New) In a method of making a seasoning, the improvement comprising producing L-glutamic acid according to the method according to claim 3.

23. (New) In a method of making a seasoning, the improvement comprising producing L-glutamic acid according to the method according to claim 6.

24. (New) In a method of making a seasoning, the improvement comprising producing L-glutamic acid according to the method according to claim 7.

25. (New) In a method of making a seasoning, the improvement comprising producing L-glutamic acid according to the method according to claim 17.

26. (New) In a method of making a seasoning, the improvement comprising producing L-glutamic acid according to the method according to claim 18.

27. (New) In a method of making a seasoning, the improvement comprising producing L-glutamic acid according to the method according to claim 19.
